## SENATE BILL NO. 510

## 102ND GENERAL ASSEMBLY

INTRODUCED BY SENATOR RAZER.

1858S.01I KRISTINA MARTIN, Secretary

## **AN ACT**

To repeal section 338.730, RSMo, and to enact in lieu thereof one new section relating to HIV preexposure prophylaxis.

Be it enacted by the General Assembly of the State of Missouri, as follows:

- Section A. Section 338.730, RSMo, is repealed and one new
- 2 section enacted in lieu thereof, to be known as section 338.730,
- 3 to read as follows:
  - 338.730. 1. Notwithstanding any other law to the
- 2 contrary, a pharmacist may dispense HIV postexposure
- 3 prophylaxis and preexposure prophylaxis in accordance with
- 4 this section. Such prophylaxis shall be dispensed only if
- 5 the pharmacist follows a written protocol authorized by a
- 6 licensed physician. A pharmacist shall not dispense more
- 7 than a thirty-day supply of preexposure prophylaxis to a
- 8 patient more than once every two years under this section;
- 9 provided, however, that a pharmacist may dispense more than
- 10 a thirty-day supply to a patient if directed by a
- 11 prescription.
- 12 2. For purposes of this section[, "postexposure
- prophylaxis" shall mean any drug approved by the Food and
- 14 Drug Administration that meets the same clinical eligibility
- 15 recommendations provided in CDC guidelines.
- 16 3. For purposes of this section, "CDC quidelines"
- 17 shall mean the current HIV guidelines published by the
- 18 federal Centers for Disease Control and Prevention] the
- 19 following terms shall mean:

EXPLANATION-Matter enclosed in bold-faced brackets [thus] in this bill is not enacted and is intended to be omitted in the law.

SB 510 2

- 20 (1) "CDC guidelines", the current HIV guidelines
  21 published by the federal Centers for Disease Control and
  22 Prevention;
- (2) "Preexposure prophylaxis", any drug approved by
  the Food and Drug Administration that meets the same
  clinical eligibility recommendations provided in CDC
  quidelines;
- 27 (3) "Postexposure prophylaxis", any drug approved by
  28 the Food and Drug Administration that meets the same
  29 clinical eligibility recommendations provided in CDC
  30 guidelines.
- [4.] 3. The state board of registration for the healing arts [and] may advise the state board of pharmacy [shall jointly] on the [promulgate] promulgation of rules and regulations for the administration of this section.

  [Neither board shall separately promulgate rules governing a pharmacist's authority to dispense HIV postexposure prophylaxis under this section.
- 5.] 4. Any rule or portion of a rule, as that term is 38 defined in section 536.010, that is created under the 39 authority delegated in this section shall become effective 40 only if it complies with and is subject to all of the 41 provisions of chapter 536 and, if applicable, section 42 43 536.028. This section and chapter 536 are nonseverable and if any of the powers vested with the general assembly 44 45 pursuant to chapter 536 to review, to delay the effective 46 date, or to disapprove and annul a rule are subsequently held unconstitutional, then the grant of rulemaking 47 48 authority and any rule proposed or adopted after August 28, 49 2021, shall be invalid and void.

**√**